

This listing of claims will replace all prior versions, and listings, of claims in the application:

### LISTING OF CLAIMS

1. (Currently Amended) An ophthalmic contact lens solution comprising:  
0.001 to 10 percent by weight ethoxylated glyceride; and  
0.001 to 2 weight percent of a physiologically acceptable buffer adjusted so the pH of solution is between 6.5 and 7.8 and the balance water and wherein said solution does not contain an iodophor.
2. (Currently Amended) An ophthalmic contact lens solution comprising:  
0.001 to 10 percent by weight ethoxylated glyceride; and  
0.001 to 2 weight percent of a physiologically acceptable tonicity agent adjusted so the solution is isotonic between 200 and 400 mOsm and wherein said solution does not contain an iodophor.
3. (Currently Amended) An ophthalmic solution comprising;  
0.001 to 10 percent by weight ethoxylated glyceride; and  
0.00001 to 0.1 weight percent of a preservative agent that do not use an iodophor.
4. (Currently Amended) The solution of claim 1 which further comprises:  
0.01 to 2 weight percent of a physiologically acceptable tonicity agent adjusted so the solution is isotonic between 200 and 400 mOsm.
5. (Original) The solution of claim 4 that further comprises 0.00001 to 0.1 weight percent of a preservative.
6. (Currently Amended) The solution of claim 1 wherein the ethoxylated glyceride is chosen from the group of compounds consisting of Polyoxyl 40 hydrogenated castor oil (Cremophor RH 40), polyoxyl 60 hydrogenated castor oil (Cremophor

RH 60), PEG-30 Castor Oil (Incrocas 30), PEG-35 Castor Oil (Cremophor EL, Incrocas 35), or PEG-40 Castor Oil (Cremophor EL, Incrocas), Cremophor EL®, Emulphor EL®, glycerol polyethyleneglycol ~~ricinoleate~~ ricinoleate, ~~glycerol~~ glycerol polyethyleneglycol oxystearate, polyethoxylated hydrogenated castor oil, [[or]] and polyethoxylated vegetable oil.

7. (Currently Amended) The solution of claim 1 wherein the buffer is selected from the group consisting of organic amines, organic carboxylic acids, amphoterics, phosphates, [[or]] and borates.
8. (Original) Method for rendering a contact lens wettable by contacting the surface of said lens with an aqueous solution comprising from .001 to about 10 percent by weight of an ethoxylated glyceride.
9. (Original) The method of claim 8 wherein the ethoxylated glyceride is polyoxyl 40 hydrogenated castor oil.
10. (Currently Amended) The method of claim [[7]] 8 wherein said ethoxylated glyceride is polyoxyl 60 hydrogenated castor oil.
11. (Currently Amended) The method of claim [[7]] 8 wherein said ethoxylated glyceride is ~~polyoxyl-40 hydrogenated~~ PEG-40 castor oil.
12. (Currently Amended) The method of claim [[7]] 8 wherein said ethoxylated glyceride is polyoxyl 35 castor oil.
13. (Currently Amended) The method of claim [[7]] 8 wherein the aqueous solution further comprises the buffer bis(2-hydroxyethyl)iminotris(hydroxymethyl)methane (Bis-Tris) and its salts.

14. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the 1,2-bis[tris(hydroxymethyl)-methylamino]propane (Bis-Tris Propane) and its salts.
15. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the N-tris(hydroxymethyl) methyl glycine (Tricine) and its salts.
16. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the N,N-bis(2-hydroxyethyl)-glycine (Bicine) and its salts.
17. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the betaine and its salts.
18. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer phosphate and its salts
19. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is borate and its salts
20. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the is citrate and its salts
21. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises is TRIS and its salts
22. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is 2-amino-2-methyl-1,3-propanediol and its salts
23. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is triisopropanolamine and its salts

24. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is carnitine and its salts
25. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is dimethyl glutamate and its salts
26. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is creatine and its salts
27. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is diethanolamine and its salts
28. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is diisopropylamine and its salts
29. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is triethanolamine and its salts
30. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is triethylamine and its salts
31. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is dimethyl aspartic acid and its salts
32. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is imidazole and its salts
33. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is histidine and its salts

34. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is methyl aspartate and its salts
35. (Withdrawn) The method of claim 7 wherein the aqueous solution further comprises the buffer is Tris(hydroxymethyl)aminomethane (Tromethamine, TRIS) and its salts
36. (Currently Amended) A contact lens product comprising:  
A contact lens:  
A sealable container; and  
An effective amount of an ophthalmic lens solution comprising:  
0.001 to 10 percent by weight ethoxylated glyceride;  
0.01 to 2 weight percent of a physiologically acceptable buffer  
adjusted so the pH of solution is between 6.5 and 7.8 and the  
balance water and wherein said solution does not contain an  
iodophor.
37. (Withdrawn) The method of claim 7 wherein the buffer is glycine and its salts
38. (Withdrawn) The method of claim 7 wherein the buffer is lysine and its salts
39. (Withdrawn) The method of claim 7 wherein the buffer is histidine and its salts.,